

GenCore version 5.1.6 (c) 1993 - 2005 Compugen Ltd.		1 nucleic - nucleic search, using SW model		Sequence 11, Appli	
Run on: November 9, 2005, 18:15:28 ; Search time 497 Seconds (without alignments)		366.068 Million cell updates/sec		Sequence 10, Appli	
Title: US-09-937-057-9		Sequence 9, Appli		Sequence 11, Appli	
Perfect score: 22		Sequence 8, Appli		Sequence 7, Appli	
Sequence: 1 tgactgtgaacgttatagatga 22		Sequence 6, Appli		Sequence 5, Appli	
Scoring table: IDENTITY_NUC		Sequence 4, Appli		Sequence 3, Appli	
Gapop 10_0 , Gapext 1.0		Sequence 2, Appli		Sequence 1, Appli	
Searched: 9794730 secs, 413490567 residues		Sequence 4, Appli		Sequence 9, Appli	
Total number of hits satisfying chosen parameters: 11332426		Sequence 12, Appli		Sequence 11, Appli	
Minimum DB seq length: 0		Sequence 10, Appli		Sequence 9, Appli	
Maximum DB seq length: 100?		Sequence 1, Appli		Sequence 0, Appli	
Post-processing: Minimum Match 0%		Sequence 4, Appli		Sequence 3, Appli	
Listing First 45 summaries		Sequence 10, Appli		Sequence 9, Appli	
Database :		Sequence 11, Appli		Sequence 8, Appli	
Published Applications Nbr.:		Sequence 7, Appli		Sequence 6, Appli	
1: /cgn2_6/ptodata/2/pubpna/us07_PUBCOMB.seq*		Sequence 5, Appli		Sequence 4, Appli	
2: /cgn2_6/ptodata/2/pubpna/pct_NEW_PUB.seq*		Sequence 3, Appli		Sequence 2, Appli	
3: /cgn2_6/ptodata/2/pubpna/us06_NEW_PUB.seq*		Sequence 1, Appli		Sequence 0, Appli	
4: /cgn2_6/ptodata/2/pubpna/us06_PUBCOMB.seq*		Sequence 11, Appli		Sequence 10, Appli	
5: /cgn2_6/ptodata/2/pubpna/us07_NEW_PUB.seq*		Sequence 9, Appli		Sequence 8, Appli	
6: /cgn2_6/ptodata/2/pubpna/pctus_PUBCOMB.seq*		Sequence 7, Appli		Sequence 6, Appli	
7: /cgn2_6/ptodata/2/pubpna/us08_NEW_PUB.seq*		Sequence 5, Appli		Sequence 4, Appli	
8: /cgn2_6/ptodata/2/pubpna/us08_PUBCOMB.seq*		Sequence 3, Appli		Sequence 2, Appli	
9: /cgn2_6/ptodata/2/pubpna/us09_PUBCOMB.seq*		Sequence 1, Appli		Sequence 0, Appli	
10: /cgn2_6/ptodata/2/pubpna/us09B_PUBCOMB.seq*		Sequence 4, Appli		Sequence 3, Appli	
11: /cgn2_6/ptodata/2/pubpna/us09C_PUBCOMB.seq*		Sequence 11, Appli		Sequence 10, Appli	
12: /cgn2_6/ptodata/2/pubpna/us09_NEW_PUB.seq*		Sequence 9, Appli		Sequence 8, Appli	
13: /cgn2_6/ptodata/2/pubpna/us10A_PUBCOMB.seq*		Sequence 7, Appli		Sequence 6, Appli	
14: /cgn2_6/ptodata/2/pubpna/us10B_PUBCOMB.seq*		Sequence 5, Appli		Sequence 4, Appli	
15: /cgn2_6/ptodata/2/pubpna/us10C_PUBCOMB.seq*		Sequence 3, Appli		Sequence 2, Appli	
16: /cgn2_6/ptodata/2/pubpna/us10D_PUBCOMB.seq*		Sequence 1, Appli		Sequence 0, Appli	
17: /cgn2_6/ptodata/2/pubpna/us10E_PUBCOMB.seq*		Sequence 11, Appli		Sequence 10, Appli	
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21: /cgn2_6/ptodata/2/pubpna/us10I_PUBCOMB.seq*		Sequence 3, Appli		Sequence 2, Appli	
22: /cgn2_6/ptodata/2/pubpna/us10J_PUBCOMB.seq*		Sequence 1, Appli		Sequence 0, Appli	
23: /cgn2_6/ptodata/2/pubpna/us10K_PUBCOMB.seq*		Sequence 11, Appli		Sequence 10, Appli	
24: /cgn2_6/ptodata/2/pubpna/us11A_NEW_PUB.seq*		Sequence 9, Appli		Sequence 8, Appli	
25: /cgn2_6/ptodata/2/pubpna/us11A_PUBCOMB.seq*		Sequence 7, Appli		Sequence 6, Appli	
26: /cgn2_6/ptodata/2/pubpna/us11B_NEW_PUB.seq*		Sequence 5, Appli		Sequence 4, Appli	
27: /cgn2_6/ptodata/2/pubpna/us11B_PUBCOMB.seq*		Sequence 3, Appli		Sequence 2, Appli	
28: /cgn2_6/ptodata/2/pubpna/us11C_PUBCOMB.seq*		Sequence 1, Appli		Sequence 0, Appli	
Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.		SUMMARIES		SUMMARIES	
Result No. Score Match Length DB ID		Query Match		Query Match	
1 22 100.0 22 9 US-09-957-881-9		Best Local Similarity		Best Local Similarity	
2 20.4 92.7 22 9 US-09-791-500-5		Matches 22;		Matches 22;	
3 20.4 92.7 22 9 US-09-791-500-6		Conservative		Conservative	
4 20.4 92.7 22 9 US-09-770-943-2		0; Mismatches		0; Mismatches	
5 20.4 92.7 22 9 US-09-848-986-11		Indels 0;		Indels 0;	
6 20.4 92.7 22 9 US-10-233-121A-10		Gaps 0;		Gaps 0;	
7 20.4 92.7 22 9 US-10-219-143-5		Sequence 11, Appli		Sequence 11, Appli	
8 20.4 92.7 22 9 US-10-219-143-6		Sequence 10, Appli		Sequence 10, Appli	
9 20.4 92.7 22 9 US-10-212-151-5		Sequence 9, Appli		Sequence 9, Appli	
10 20.4 92.7 22 9 US-10-412-151-6		Sequence 8, Appli		Sequence 8, Appli	
11 20.4 92.7 22 9 US-10-233-121A-11		Sequence 7, Appli		Sequence 7, Appli	
12 19.4 88.2 22 9 US-09-802-685-1		Sequence 6, Appli		Sequence 6, Appli	
13 18.8 85.5 22 9 US-09-802-685-12		Sequence 5, Appli		Sequence 5, Appli	
14 18.8 85.5 22 9 US-09-802-685-14		Sequence 4, Appli		Sequence 4, Appli	
15 18.8 85.5 22 9 US-09-802-685-15		Sequence 3, Appli		Sequence 3, Appli	
16 18.8 85.5 22 9 US-09-802-685-1		Sequence 2, Appli		Sequence 2, Appli	
17 18.8 85.5 22 9 US-09-802-685-17		Sequence 1, Appli		Sequence 1, Appli	
18 18.8 85.5 22 9 US-09-802-685-19		Sequence 0, Appli		Sequence 0, Appli	
19 18.8 85.5 22 9 US-09-802-685-20		Sequence 1, Appli		Sequence 1, Appli	
20 18.8 85.5 22 9 US-09-802-685-21		Sequence 0, Appli		Sequence 0, Appli	
21 18.8 85.5 22 9 US-09-791-500-1		Sequence 1, Appli		Sequence 1, Appli	
22 18.8 85.5 22 9 US-09-791-500-2		Sequence 0, Appli		Sequence 0, Appli	
23 18.8 85.5 22 9 US-09-791-500-3		Sequence 1, Appli		Sequence 1, Appli	
24 18.8 85.5 22 9 US-09-791-500-4		Sequence 0, Appli		Sequence 0, Appli	
25 18.8 85.5 22 9 US-09-791-500-5		Sequence 1, Appli		Sequence 1, Appli	
26 18.8 85.5 22 9 US-09-791-500-6		Sequence 0, Appli		Sequence 0, Appli	
27 18.8 85.5 22 9 US-09-791-500-7		Sequence 1, Appli		Sequence 1, Appli	
28 18.8 85.5 22 9 US-09-791-500-8		Sequence 0, Appli		Sequence 0, Appli	
29 18.8 85.5 22 9 US-09-791-500-9		Sequence 1, Appli		Sequence 1, Appli	
30 18.8 85.5 22 9 US-09-770-943-1		Sequence 0, Appli		Sequence 0, Appli	
31 18.8 85.5 22 9 US-09-802-370-1		Sequence 1, Appli		Sequence 1, Appli	
32 18.8 85.5 22 9 US-09-802-370-4		Sequence 0, Appli		Sequence 0, Appli	
33 18.8 85.5 22 9 US-09-802-445-1		Sequence 1, Appli		Sequence 1, Appli	
34 18.8 85.5 22 9 US-09-802-445-2		Sequence 0, Appli		Sequence 0, Appli	
35 18.8 85.5 22 9 US-09-802-445-3		Sequence 1, Appli		Sequence 1, Appli	
36 18.8 85.5 22 9 US-09-820-484-3		Sequence 0, Appli		Sequence 0, Appli	
37 18.8 85.5 22 9 US-09-820-484-4		Sequence 1, Appli		Sequence 1, Appli	
38 18.8 85.5 22 9 US-09-820-484-5		Sequence 0, Appli		Sequence 0, Appli	
39 18.8 85.5 22 9 US-09-828-505-1		Sequence 1, Appli		Sequence 1, Appli	
40 18.8 85.5 22 9 US-09-828-505-2		Sequence 0, Appli		Sequence 0, Appli	
41 18.8 85.5 22 9 US-09-967-881-1		Sequence 1, Appli		Sequence 1, Appli	
42 18.8 85.5 22 9 US-09-967-881-2		Sequence 0, Appli		Sequence 0, Appli	
43 18.8 85.5 22 9 US-09-967-881-3		Sequence 1, Appli		Sequence 1, Appli	
44 18.8 85.5 22 9 US-09-927-422A-1		Sequence 0, Appli		Sequence 0, Appli	
45 18.8 85.5 22 9 US-09-927-422A-4		Sequence 1, Appli		Sequence 1, Appli	
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; Sequence 9, Application US-09967881		; Sequence 9, Application US-09967881		; Sequence 9, Application US-09967881	
; Publication No. US200201921841		; Publication No. US200201921841		; Publication No. US200201921841	
; GENERAL INFORMATION:		; GENERAL INFORMATION:		; GENERAL INFORMATION:	
; APPLICANT: Assistance Publique - Hopitaux de Paris		; APPLICANT: Assistance Publique - Hopitaux de Paris		; APPLICANT: Assistance Publique - Hopitaux de Paris	
; INVENTION: Use of Stabilised Oligonucleotides for Preparing A Medicament		; INVENTION: Use of Stabilised Oligonucleotides for Preparing A Medicament		; INVENTION: Use of Stabilised Oligonucleotides for Preparing A Medicament	
; TITLES OF INVENTION: Antitumor Activity		; TITLES OF INVENTION: Antitumor Activity		; TITLES OF INVENTION: Antitumor Activity	
; FILE REFERENCES: 267/246 US		; FILE REFERENCES: 267/246 US		; FILE REFERENCES: 267/246 US	
; CURRENT APPLICATION NUMBER: US/09/967,881		; CURRENT APPLICATION NUMBER: US/09/967,881		; CURRENT APPLICATION NUMBER: US/09/967,881	
; CURRENT FILING DATE: 2001-09-28		; CURRENT FILING DATE: 2001-09-28		; CURRENT FILING DATE: 2001-09-28	
; SOFTWARE: Patentin version 3.1		; SOFTWARE: Patentin version 3.1		; SOFTWARE: Patentin version 3.1	
; SEQ ID NO: 9		; SEQ ID NO: 9		; SEQ ID NO: 9	
; LENGTH: 22		; LENGTH: 22		; LENGTH: 22	
; TYPE: DNA		; TYPE: DNA		; TYPE: DNA	
; ORGANISM: Artificial sequence		; ORGANISM: Artificial sequence		; ORGANISM: Artificial sequence	
; FEATURE:		; FEATURE:		; FEATURE:	
; OTHER INFORMATION: Oligodeoxynucleotide		; OTHER INFORMATION: Oligodeoxynucleotide		; OTHER INFORMATION: Oligodeoxynucleotide	
; US-09-967-881-9		; US-09-967-881-9		; US-09-967-881-9	

RESULT 7
 TITLE OF INVENTION: Methods of Use Thereof
 FILE REFERENCE: 06510168051
 CURRENT APPLICATION NUMBER: US/09/848,986
 CURRENT FILING DATE: 2001-05-03
 PRIORITY NUMBER: US 60/262321
 PRIORITY FILING DATE: 2001-01-17
 PRIORITY APPLICATION NUMBER: US 60/202,274
 PRIORITY FILING DATE: 2000-05-05
 NUMBER OF SEQ ID NOS: 21
 SEQ ID NO: 11
 LENGTH: 22
 TYPE: DNA
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: methylated ISS-ODN
 NAME/KEY: modified base
 LOCATION: (11)...(11)
 OTHER INFORMATION: m5C
 US-09-848-986-11

Query Match 92.7%; Score 20.4; DB 10; Length 22;
 Best Local Similarity 95.5%; Pred. No. 9.9;
 Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 TGACCTGAACTTATAGATGA 22
 Db 1 TGACCTGAACTTATAGATGA 22

RESULT 8
 US-10-233-121A-10
 Sequence 10, Application US/10233121A
 Publication No. US20030125284A1
 GENERAL INFORMATION:
 APPLICANT: RAZ, EYAL
 APPLICANT: LOIS, AUGUSTO
 APPLICANT: TAKABAYASHI, KENJI
 TITLE OF INVENTION: METHODS OF USE THEREOF
 FILE REFERENCE: UCAL-16DIV
 CURRENT APPLICATION NUMBER: US/10/233,121A
 CURRENT FILING DATE: 2003-03-11
 PRIORITY NUMBER: US 09/848,986
 PRIORITY FILING DATE: 2001-05-04
 PRIORITY APPLICATION NUMBER: US 60/202,274
 PRIORITY FILING DATE: 2000-05-05
 PRIORITY APPLICATION NUMBER: US 60/262,321
 PRIORITY FILING DATE: 2001-01-17
 NUMBER OF SEQ ID NOS: 21
 SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO: 10
 LENGTH: 22
 TYPE: DNA
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: phosphodiester or phosphorothioate oligonucleotide

Query Match 92.7%; Score 20.4; DB 16; Length 22;
 Best Local Similarity 95.5%; Pred. No. 9.9;
 Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 1 TGACCTGAACTTATAGATGA 22
 Db 1 TGACCTGAACTTATAGATGA 22

RESULT 9
 US-10-219-143-6
 Sequence 6, Application US/10219143
 Publication No. US20030130217A1
 GENERAL INFORMATION:
 APPLICANT: RAZ, EYAL
 APPLICANT: Rachmilewitz, Daniel
 TITLE OF INVENTION: Disease and Other Forms of Gastrointestinal Inflammation.
 FILE REFERENCE: 6510-00US1
 CURRENT APPLICATION NUMBER: US/10/219,143
 CURRENT FILING DATE: 2002-08-13
 PRIORITY NUMBER: US/09/791,500
 PRIORITY FILING DATE: 2001-04-22
 NUMBER OF SEQ ID NOS: 39
 SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO: 5
 LENGTH: 22
 TYPE: DNA
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: synthetic polynucleotide sequence
 US-10-219-143-5

RESULT 10
 US-10-412-151-5
 Sequence 5, Application US/10412151
 Publication No. US20030176389A1
 GENERAL INFORMATION:
 APPLICANT: RAZ, EYAL
 APPLICANT: Rachmilewitz, Daniel
 TITLE OF INVENTION: Method for Treating Inflammatory Bowel
 FILE REFERENCE: UCAL-202CON
 CURRENT APPLICATION NUMBER: US/10/412,151
 CURRENT FILING DATE: 2003-04-11

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; PRIOR APPLICATION NUMBER: 09/791,500
; PRIOR FILING DATE: 2001-02-12
; PRIOR APPLICATION NUMBER: 60/184,256
; PRIOR FILING DATE: 2000-02-23
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 5
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE: OTHER INFORMATION synthetic polynucleotide sequence
; OTHER INFORMATION: synthetic polynucleotide sequence
US-10-412-151-5

Query Match 92.7%; Score 20.4; DB 17; Length 22;
Best Local Similarity 95.5%; Pred. No. 9, 9;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 TGACTGAACTGTTAGATGA 22
Db 1 TGACTGAACTGTTAGATGA 22

RESULT 11
US-10-412-151-6
; Sequence 6, Application US/10412151
; Publication No. US2003017638941
; GENERAL INFORMATION
; APPLICANT: Raz, Eyal
; APPLICANT: Rachmlevitz, Daniel
; TITLE OF INVENTION: Method for Treating Inflammatory Bowel Disease and Other Forms of Gastrointestinal Inflammation.
; FILE REFERENCE: UCAL-202CON
; CURRENT APPLICATION NUMBER: US/10/412,151
; CURRENT FILING DATE: 2003-04-11
; PRIOR APPLICATION NUMBER: 09/791,500
; PRIOR FILING DATE: 2001-02-22
; PRIOR APPLICATION NUMBER: 60/184,256
; PRIOR FILING DATE: 2000-02-13
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 6
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE: OTHER INFORMATION synthetic polynucleotide sequence
; OTHER INFORMATION: synthetic polynucleotide sequence
US-10-412-151-6

Query Match 92.7%; Score 20.4; DB 17; Length 22;
Best Local Similarity 95.5%; Pred. No. 9, 9;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 TGACTGAACTGTTAGATGA 22
Db 1 TGACTGAACTGTTAGATGA 22

RESULT 12
US-10-233-121A-11
; Sequence 11, Application US/10233121A
; Publication No. US20030125284A1
; GENERAL INFORMATION
; APPLICANT: RAZ, EYAL
; APPLICANT: LOIS, AUGUSTO
; APPLICANT: TAKABAYASHI, KENJI
; TITLE OF INVENTION: AGENTS THAT MODULATE DNA-PK ACTIVITY AND METHODS OF USE THEREOF
; FILE REFERENCE: UCAL-158DIV
; CURRENT APPLICATION NUMBER: US/10/233,121A
; CURRENT FILING DATE: 2003-03-11
; PRIOR APPLICATION NUMBER: US 09/848,986
; PRIOR FILING DATE: 2001-05-04

Query Match 88.2%; Score 19.4; DB 16; Length 22;
Best Local Similarity 90.9%; Pred. No. 30;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 1 TGACTGAACTTATAGATGA 22
Db 1 TGACTGAACTTATAGATGA 22

RESULT 13
US-09-802-686-1
; Sequence 1, Application US/09802686
; Patent No. US200104697A1
; GENERAL INFORMATION
; APPLICANT: Dynavax Technologies Corporation
; APPLICANT: Van Nest, Gary
; TITLE OF INVENTION: METHODS OF PREVENTING AND TREATING RESPIRATORY VIRAL INFECTION USING IMMUNOMODULATORY POLYNUCLEOTIDE SEQUENCES
; FILE REFERENCE: 377882000900
; CURRENT APPLICATION NUMBER: US/09/802,686
; CURRENT FILING DATE: 2001-03-09
; PRIOR APPLICATION NUMBER: 60/188,583
; PRIOR FILING DATE: 2000-03-10
; NUMBER OF SEQ ID NOS: 10
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE: OTHER INFORMATION: Polynucleotide containing CG
US-09-802-686-1

Query Match 85.5%; Score 18.8; DB 9; Length 22;
Best Local Similarity 90.9%; Pred. No. 57;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 1 TGACTGAACTTATAGATGA 22
Db 1 TGACTGAACTTATAGATGA 22

RESULT 14
US-09-802-686-4
; Sequence 4, Application US/09802686
; Patent No. US200104697A1
; GENERAL INFORMATION
; APPLICANT: Dynavax Technologies Corporation
; APPLICANT: Van Nest, Gary
; TITLE OF INVENTION: METHODS OF PREVENTING AND TREATING RESPIRATORY VIRAL INFECTION USING IMMUNOMODULATORY POLYNUCLEOTIDE SEQUENCES
; FILE REFERENCE: 377882000900
; CURRENT APPLICATION NUMBER: US/09/802,686
; CURRENT FILING DATE: 2001-03-09

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PRIOR APPLICATION NUMBER: 60/188, 583
PRIOR FILING DATE: 2000-03-10
NUMBER OF SEQ ID NOS: 10
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO: 4
LENGTH: 22
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Polynucleotide containing CG
us-09-802-686-4

Query Match 85.5%; Score 18.8; DB 9; Length 22;
Best Local Similarity 90.9%; Pred. No. 57;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 1 TGACTGTAACGTTAGATGA 22
1 TGACTGTAACGTTCCAGATGA 22

RESULT 15

us-09-802-686-9
Sequence 9, Application US/09802686
Patent No. US20010046967A1
GENERAL INFORMATION:
APPLICANT: Dynavax Technologies Corporation
APPLICANT: Van Ness, Gary
TITLE OF INVENTION: METHODS OF PREVENTING AND TREATING
TITLE OF INVENTION: RESPIRATORY VIRAL INFECTION USING IMMUNOMODULATORY
TITLE OF INVENTION: POLYNUCLEOTIDE SEQUENCES
FILE REFERENCE: 377882000900
CURRENT APPLICATION NUMBER: US/09/802, 686
CURRENT FILING DATE: 2001-03-09
PRIOR APPLICATION NUMBER: 60/188, 583
PRIOR FILING DATE: 2000-03-10
NUMBER OF SEQ ID NOS: 10
SOFTWARE: FastSEQ for Windows Version 4.0
SEQ ID NO 9
LENGTH: 22
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Polynucleotide not containing CG
us-09-802-686-9

Query Match 85.5%; Score 18.8; DB 9; Length 22;
Best Local Similarity 90.9%; Pred. No. 57;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy ||||| ||||| ||||| ||||| ||||| |||||
Db 1 TGACTGTAACGTTAGATGA 22
1 TGACTGTAACGTTCCAGATGA 22

Search completed: November 9, 2005, 19:28:46
Job time : 504 secs

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/organism="Mus musculus"
 /mol type="mRNA"
 /strain="29Sv/Ev"
 /db_xref="taxon:10090"
 /clone="OSM36752"
 /cell type="embryonic stem cell"
 /clone_lib="Mus musculus 129Sv/Ev"

ORIGIN

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 Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Db 49 TGACCGGAACTGATAGAGGA 70

RESULT 8
 AIG62714/c

LOCUS AIG62714 58 bp mRNA linear EST 10-MAY-1999
 DEFINITION va8bc01.x1 Soares mouse 3NME12.5 Mus musculus cDNA clone IMAGE:746496 3' Similar to TR:088760 088760 AF-9 PROTEIN. ; mRNA sequence.

ACCESSION AIG62714
 VERSION 1
 KEYWORDS EST.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 58)
 AUTHORS Marra, M., Hillier, L., Kubaca, T., Martin, J., Beck, C., Wylie, T., Underwood, K., Stephre, M., Theising, B., Allen, M., Bowers, Y., Person, B., Swaller, T., Gibbons, M., Pape, D., Harvey, N., Schurk, R., Ritter, E., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and Wilson, R.

TITLE The WashU-NCI Mouse EST Project 1999
 JOURNAL Other ESTs: va88c01.y1
 COMMENT Contact: Marra M/WashU-NCI Mouse EST Project 1999 Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA Tel: 314 286 1800 Fax: 314 286 1810 Email: mouseest@watson.wustl.edu
 This clone is available royalty-free through LInI; contact the IMAGE Consortium (info@image.lnl.gov) for further information.

MGFI-455480

This clone was previously sequenced on the 5' end only, this new data is from the 3' end. Possible reversed clone: similarity on wrong strand

High quality sequence stop: 1.

FEATURES SOURCE
 1. .58
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 /mol type="mRNA"
 /strain="29Sv/Ev"
 /db_xref="taxon:10090"
 /clone IMAGE:746496"
 /sex="unknown"
 /tissue type="fetus"
 /dev_stage="12.5dpc total fetus"
 /lab host="DH10B"
 /clone lib="Soares mouse 3NME12.5"
 /note="Organ: whole fetus; Vector: PT7T3D-Pac (Pharmacia) with a modified polylinker; Site: 1 - Not I - Oligo(dT) primer 1st strand cDNA was primed with a Not I - Oligo(dT) primer 3', 1, on total mouse RNA (provided by Minoru Ko, Wayne State Univ.); double stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified PT7T3 vector.

ORIGIN

Query Match 2 GACTGTGAACCTATAGATGA 22
 Best Local Similarity 81.0%; Score 14.6; DB 1; Length 58;
 Matches 17; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Db 29 GATTGGACCTATAGAGGA 9

RESULT 9
 BH908898

LOCUS BH908898 58 bp DNA linear GSS 04-SEP-2002

DEFINITION SALK_051118.17.90.x Arabidopsis thaliana TDNA insertion lines

Arabidopsis thaliana genomic clone SALK_051118.17.90.x, genomic survey sequence.

BH908898

ACCESSION BH908898.1 GI:22721831

KEYWORDS GSS.

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM Arabidopsis thaliana

Bukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicots; eudicots; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE 1 (bases 1 to 58)
 Alonso J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R., Gastrinab, C., Jester, A., Kernes, M., Kim, C.J., Parker, H., Prednis, L., Shinn, P., Zimmerman, J., and Ecker, J.R., Unpublished (2001)

AUTHORS Alonso J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R., Gastrinab, C., Jester, A., Kernes, M., Kim, C.J., Parker, H., Prednis, L., Shinn, P., Zimmerman, J., and Ecker, J.R., Unpublished (2001)

COMMENT Contact: Joseph R. Ecker
 Salk Institute Genomic Analysis Laboratory (SIGnAL)
 The Salk Institute for Biological Studies
 10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
 Tel: 858 453 4100 x1752
 Fax: 858 558 6379
 Email: ecker@salk.edu

This is single pass sequence recovered from the left border of TDNA. This sequence lies within an annotated intron of At1g71692.

FEATURES SOURCE
 1. .58
 /organism="Arabidopsis thaliana"
 /mol type="genomic DNA"
 /ecotype="Col-0"
 /db xref="taxon:3702"
 /clone=SALK_051118.17.90.x"
 /clone lib="Arabidopsis thaliana TDNA insertion lines"
 /note="PCR was performed on Arabidopsis thaliana lines each of which contains one or more TDNA insertion elements. The resultant fragment for each line was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at http://signal.salk.edu/tDNA_protocols.html"

ORIGIN

Query Match 1 TGACTGTGAACGTTATAGATG 21
 Best Local Similarity 81.0%; Score 14.6; DB 8; Length 58;
 Matches 17; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Db 9 TTACTGTAAAGTAAATAGATG 29

RESULT 10
 CG546804/c

LOCUS CG546804 70 bp mRNA linear GSS 01-OCT-2003

DEFINITION OST146791 Mus musculus 129Sv/Ev Mus musculus cDNA clone OST146791,

ACCESSION	mRNA sequence.	FEATURES	Seq primer: T3 backward
VERSION	CG546804	source	POLYA>No. 1..80
KEYWORDS	GSS		Location/Qualifiers
SOURCE	Mus musculus (house mouse)		/organism="Phaeodactylum tricornutum"
ORGANISM	Mus musculus		/mol_type="mRNA"
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Buteleostomi; Mammalia; Eutheria; Rodentia; (bases 1 to 70)		/db_xref="taxon:2850"
AUTHORS	Zambrowicz, B.P., Abuin, A., Ramirez-Solis, R., Richter, L.J., Piggott, J.J., Beltran, D.R.H., Burton, R.C., Edwards, J., Finch, R.A., Friddle, C.J., Gupta, A., Hansen, G., Hu, Y., Huang, W., Jaing, C., Key, B.W. Jr., Kipp, P., Kohlhauff, B., Ma, Z.-Q., Markesich, D., Sparks, M.J., Spurr, D.G., Qian, N., Shaw, J., Schrick, J., Shi, Z., Zhou, Q., Person, C. and Sands, A.T.	ORIGIN	/cell_line="CCMP632"
TITLE	Wnk1 kinase deficiency lowers blood pressure in mice: a gene-trap screen to identify potential targets for therapeutic intervention		/note="Vector: Uni-Zap XR vector; Site_1: Eco RI; Site_2: Xba I"
JOURNAL	Proc. Natl. Acad. Sci. U.S.A. 100 (24), 14109-14114 (2003)		
COMMENT	OmniBank		
Lexicon Genetics Incorporated	4000 Research Forest Drive, The Woodlands, TX 77381, USA		
	Email: materials@lexgen.com		
	Gene trap sequence tag generated by 3' RACE from mouse ES cells as described in Zambrowicz et al (Nature. 1998 Apr 9;392(6676):608-11)		
	Class: Gene Trap.		
FEATURES	Location/Qualifiers		
source	1..70		
	/organism="Mus musculus"		
	/mol_type="mRNA"		
	/strain="129Sv/Ev"		
	/db_xref="taxon:10900"		
	/clone="OST146791"		
	/cell_type="embryonic stem cell"		
	/clone_lib="Mus musculus 129Sv/Ev"		
ORIGIN			
	Query Match	Query Match	Score 14.4%; DB 9; Length 70;
	Best Local Similarity	Best Local Similarity	93.8%; Pred. No. 2.2e+04;
	Matches 15;	Matches 15;	Conservative 0; Mismatches 0;
Qy	7 TGAACTGTATAGTGA	Qy	7 TGAACTGTATAGTGA
Db	67 TGAAAGTGTATAGTGA	Db	67 TGAAAGTGTATAGTGA
MEDLINE	22111123		52
PUBMED	12114555		
COMMENT	Contact: Bowler C		
	Laboratory of Molecular Plant Biology		
	Stazione Zoologica Anton Dohrn,		
	Villa Comunale, I-80121, Napoli, Italy		
	Tel: 39 081 583 3288/3221		
	Email: chris@alpha.szn.it		
	Diatom EST Database http://avesthagen.sznbowler.com		

RESULT 13			ACCESSION AZ843479	VERSION AZ843477.1	GI:13013387
BB881470	97 bp mRNA linear EST 16-SEP-2002		KEYWORDS GSS,		
LCUS fm93a10.y1	Zebrafish Research Genetics C32 Fin Danio rerio cDNA		SOURCE Mus musculus (house mouse)		
DEFINITION clone IMAGB:4468146 5' similar to TR:Q9W725 Q9W725 UNCOUPLING PROTEIN 2. ; mRNA sequence.			ORGANISM Mus musculus		
ACCESSION BB881470			Eukaryota; Metazoa; Chordata; Craniata; Chordata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.		
VERSION 1.			REFERENCE 1 (bases 1 to 48)		
KEYWORDS EST.			AUTHORS Dunn, D.; Aoyagi, A.; Barber, M.; Beacons, T.; Duval, B.; Hamm, C.; Islam, H.; Longacre, S.; Mahmoud, M.; Meenen, E.; Pedersen, T.; Reilly, M.; Rose, M.; Rose, R.; Stokes, R.; Tingey, A.; von Niederhäusern, A. and Wright, D.; Weiss, R.		
ORGANISM Danio rerio (zebrafish)			TITLE Mouse whole genome scaffolding with paired end reads from 10kb plasmid inserts		
ACCESSION BB881470.1	EST.		COMMENT Unpublished (2000)		
VERSION 1.			CONTACT Robert B. Weiss		
KEYWORDS			University of Utah Genome Center		
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Cypriniformes; Cyprinidae; Danio.			University of Utah		
REFERENCE 1 (bases 1 to 97)			Rm. 3038, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT		
AUTHORS Clark, M.; Johnson, S. L.; Lehrach, H.; Lee, R.; Li, F.; Marra, M.; Eddy, S.; Hilliker, L.; Kucaba, T.; Martin, J.; Beck, C.; Wylie, T.; Underwood, K.; Stepcik, M.; Theising, B.; Allen, M.; Bowers, Y.; Person, B.; Swaller, T.; Gibbons, M.; Pape, D.; Harvey, N.; Schurk, R.; Ritter, B.; Kohn, S.; Shin, T.; Jackson, Y.; Cardenas, M.; McCann, R.; Waterston, R. and Wilson, R.			84112, USA		
TITLE WashU Zebrafish EST Project 1998			TEL: 801 585 5606		
JOURNAL Unpublished (1998)			FAX: 801 585 7177		
COMMENT Contact: Stephen L. Johnson			EMAIL: ddunn@genetics.utah.edu		
CDNA Library Preparation: Ning Wu. CDNA Library Arrayed by: Steve Johnson. DNA Sequencing by: Washington University Genome Sequencing Center. NONE: This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info.llnl.gov) for further information.			Insert Length: 10000 Std. Error: 0.00		
CDNA Library Preparation: Ning Wu. CDNA Library Arrayed by: Steve Johnson. DNA Sequencing by: Washington University Genome Sequencing Center. NONE: This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info.llnl.gov) for further information.			Plate: 0142 row: C column: 23		
CONTACT: Stephen L. Johnson			Seq Primer: CGTGTAAAGACGCCAGT		
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA			Class: plasmid ends		
Fax: 314 286 1800			High quality sequence stop: 48.		
email: zebrafish@watson.wustl.edu			Location/Qualifiers 1 . 48		
CDNA Library Preparation: Ning Wu. CDNA Library Arrayed by: Steve Johnson. DNA Sequencing by: Washington University Genome Sequencing Center. NONE: This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info.llnl.gov) for further information.			/organism="Mus musculus"		
Trace considered overall poor quality			/mol_type="genomic DNA"		
Seq prime: T3 ET from Amersham			/strain="C57BL/6J"		
High quality sequence stop: 1.			/lab_host="E. Coli strain XL10-Gold, T1-resistant, F-		
Location/Qualifiers 1 . 97			/note="Vector: pWPA2nv; Purified genomic DNA from M. musculus C57BL/6J (male) was obtained from the Jackson Laboratory Mouse DNA Resource		
FEATURES source			(http://www.jax.org/resources/documents/dnases/). The DNA was hydrodynamically sheared by repeated passage through a 0.005 inch orifice at constant velocity. The sheared DNA was blunt end-repaired with T4 DNA polymerase and T4 polynucleotide kinase. Adaptor oligonucleotides were ligated to the blunt ends in high molar excess. The adaptor DNA was purified and size-selected for a 9.5 to 10.5 kb range using preparative agarose gel electrophoresis. Vector DNA was prepared from a derivative of pWD42 (gi 473214 gb AF10972.1), a copy-number indelible derivative of plasmid R1. The vector was ligated with adaptors complementary to the insert adaptors and purified. The sheared, adaptor-modified mouse DNA was annealed to adaptor vector DNA, and transformed into chemically-competent E. coli XL10-Gold (Stratagene) cells and selected for ampicillin resistance."		
FEATURES source			ORIGIN		
/clone lib="zebrafish Research Genetics C32 fin"			Query Match 63 6%; Score 14; DB 8; Length 48;		
/note="Vector: pRTBD-Pac with a modified polylinker; Site 1: EcoRI; Site 2: NotI; 1st strand cDNA was prepared from zebrafish(C32) fin, and was then primed with a Not I - Oligo (dT) primer. Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pRT73 vector. Library is non-normalized. Library was constructed by Ning Wu. NOTE: This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info.llnl.gov) for further information."			Best Local Similarity 77.3%; Pred. No. 3.2e+04; Matches 17; Conservative 0; Mismatches 5; Indels 0; Gaps 0;		
DEFINITION			Qy 1 TGACTGTGAACTTATAGATGA 22		
LOCUS 1M216018R	48 bp DNA linear		Db 37 TGACTGTGAACTTATAGATGA 16		
DEFINITION 2M0142C23R	Mouse 10kb plasmid UGGC1M library				
DEFINITION Clone UGGC2M0142C23	Genomic survey sequence.				
RESULT 14					
A2843479	48 bp DNA linear				
LCUS 2M0142C23R	Mus musculus genomic				
DEFINITION	Genomic survey sequence.				

ACCESSION AZ431742
 VERSION AZ431742.1
 KEYWORDS GSS
 SOURCE Mus musculus (house mouse)
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Buteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus;
 1 (bases 1 to 63)
 REFERENCE Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,
 Islam, H., Longacre, S., Mahmoud, M., Moenen, E., Pedersen, T.,
 Reilly, M., Rose, M., Rose, R., Stokes, R., Tingey, A., von
 Niederhausern, A. and Wright, D., Weiss, R.
 TITLE Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
 JOURNAL Unpublished (2000)
 COMMENT Contact: Robert B. Weiss
 University of Utah
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
 84112, USA
 Tel: 801 585 5606
 Fax: 801 585 7177
 Email: ddunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0216 row: O column: 18
 Seq primer: CACACAGAAACAGCTATGACC
 Class: Plasmid ends
 FEATURES High quality sequence stop: 63.
 source Location/Qualifiers
 1. .63
 /organism="Mus musculus"
 /mol_type="genomic DNA"
 /strain="C57BL/6J"
 /db_xref="itaxon:10090"
 /clone="UVC1M0216018"
 /sex="Male"
 /lab_host="E. Coli strain XL10-Gold, λ -resistant, P-
 /lab_name="Mouse 10 kb Plasmid UVC1M library"
 /note="Vector: PWD2nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (<http://wwwjax.org/resources/documents/dnare/>). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adapter oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adapter DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pMD42 (91|732114|9b AF123072.1), a copy number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adapter mouse DNA was annealed to
 adapter vector DNA, and transformed into
 chemically-competent *E. coli* XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

ORIGIN

	Query Match	Score	DB	Length
Best Local Matches	17;	14;	8;	63;
	Similarity	73.34;	Pred. No.	3.4e+04;
	Conservative	0;	Mismatches	5;
			Indels	0;
			Gaps	0;

Qy 1 TCACTGTGAACTTATAGATGA 22
 Db 36 TGAATGTGAATGTGAAATGA 57

Search completed: November 9, 2005, 19:18:31
 Job time : 1772 secs

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; TYPE: DNA ; ORGANISM: Artificial Sequence
; FEATURE: OTHER INFORMATION: Synthetic polynucleotide sequence
; US-09-791-500-5

Query Match 92.7%; Score 20.4; DB 4; Length 22;
Best Local Similarity 95.5%; Pred. No. 1;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
; US-09-092-314-3
; Sequence 3, Application US/09092314
; Patent No. 6225292
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; ATTORNEY: Roman, Mark
; TITLE OF INVENTION: Method For Treating Inflammatory Bowel Disease and Other Forms of Gastrointestinal Inflammation.
; FILE REFERENCE: 6510-173US1
; CURRENT APPLICATION NUMBER: US/09/791.500
; CURRENT FILING DATE: 2001-02-22
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 6
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE: OTHER INFORMATION: Synthetic polynucleotide sequence
; US-09-791-500-6

Query Match 92.7%; Score 20.4; DB 4; Length 22;
Best Local Similarity 95.5%; Pred. No. 1;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
; US-09-092-314-3
; Sequence 6, Application US/09191500
; Patent No. 6613751
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; ATTORNEY: Rachmlevitz, Daniel
; TITLE OF INVENTION: Method For Treating Inflammatory Bowel Disease and Other Forms of Gastrointestinal Inflammation.
; FILE REFERENCE: 6510-202US1
; CURRENT APPLICATION NUMBER: US/09/791.500
; CURRENT FILING DATE: 2001-02-22
; NUMBER OF SEQ ID NOS: 39
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 6
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE: OTHER INFORMATION: Synthetic polynucleotide sequence
; US-09-791-500-5

Query Match 92.7%; Score 20.4; DB 4; Length 22;
Best Local Similarity 95.5%; Pred. No. 1;
Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
; US-09-092-314-3
; Sequence 1, Application US/09092314
; Patent No. 6225292
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; ATTORNEY: Roman, Mark
; TITLE OF INVENTION: Inhibitors of DNA Immunostimulatory Sequence Activity
; FILE REFERENCE: 6510-173US1
; CURRENT APPLICATION NUMBER: US/09/092-314
; CURRENT FILING DATE: 1998-06-05
; PRIOR APPLICATION NUMBER: 60/048,794
; PRIOR FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE: OTHER INFORMATION: Oligonucleotide
; US-09-092-314-1

Query Match 85.5%; Score 18.8; DB 3; Length 22;
Best Local Similarity 90.9%; Pred. No. 6.3;
Matches 20; Conservative 0; Mismatches 0;
Indels 0; Gaps 0;
; US-09-092-314-1
; Sequence 3, Application US/09092314
; Patent No. 6225292
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; ATTORNEY: Roman, Mark
; TITLE OF INVENTION: Inhibitors of DNA Immunostimulatory Sequence Activity
; FILE REFERENCE: 6510-173US1
; CURRENT APPLICATION NUMBER: US/09/092-314
; CURRENT FILING DATE: 1998-06-05
; PRIOR APPLICATION NUMBER: 60/048,794
; PRIOR FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE: OTHER INFORMATION: Oligonucleotide
; US-09-092-314-1

Query Match 85.5%; Score 18.8; DB 3; Length 22;
Best Local Similarity 90.9%; Pred. No. 6.3;
Matches 20; Conservative 0; Mismatches 0;
Indels 0; Gaps 0;
; US-09-092-314-1
; Sequence 1, Application US/09092314
; Patent No. 6225292
; GENERAL INFORMATION:
; APPLICANT: Raz, Eyal
; ATTORNEY: Roman, Mark
; TITLE OF INVENTION: Inhibitors of DNA Immunostimulatory Sequence Activity
; FILE REFERENCE: 6510-173US1
; CURRENT APPLICATION NUMBER: US/09/092-314
; CURRENT FILING DATE: 1998-06-05
; PRIOR APPLICATION NUMBER: 60/048,794
; PRIOR FILING DATE: 1997-06-06
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 22
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE: OTHER INFORMATION: Oligonucleotide
; US-09-092-314-1

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RESULT 7
 US-09-235-742-19
 ; Sequence 1, Application US/09235742
 ; Patent No. 6498148
 ; GENERAL INFORMATION:
 ; APPLICANT: Raz, Eyal
 ; TITLE OF INVENTION: Immunization-Free Methods for Treating
 ; TITLE OF INVENTION: Antigen-Stimulated Inflammation in a Mammalian Host and
 ; TITLE OF INVENTION: Shifting the Host's Antigen Immune Responsiveness to a TH1
 ; FILE REFERENCE: 6510-170CON4
 ; CURRENT APPLICATION NUMBER: US/09/235,742
 ; CURRENT FILING DATE: 1999-01-21
 ; EARLIER APPLICATION NUMBER: 08/9227,120
 ; EARLIER FILING DATE: 1997-09-05
 ; EARLIER APPLICATION NUMBER: 08/593,554
 ; EARLIER FILING DATE: 1996-01-30
 ; EARLIER APPLICATION NUMBER: 08/725,968
 ; EARLIER FILING DATE: 1996-10-04
 ; EARLIER APPLICATION NUMBER: 60/028,118
 ; EARLIER FILING DATE: 1996-10-11
 ; NUMBER OF SEQ ID NOS: 20
 ; SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 19
 LENGTH: 22
 TYPE: DNA
 ORGANISM: Artificial Sequence
 FEATURE: Recombinant or Synthetic Sequence
 OTHER INFORMATION: Recombinant or Synthetic Sequence
 US-09-235-742-19

Query Match 85.5%; Score 18.8; DB 4; Length 22;
 Best Local Similarity 90.9%; Pred. No. 6.3;
 Matches 20; Conservative 0; Mismatches 2; Indels 0;
 Gaps 0;
 Db 1 TGACGTGAACTTATAGATGA 22
 1 ||||| ||||| ||||| |||||

RESULT 8
 US-09-347-343-32
 ; Sequence 32, Application US/09347343A
 ; Patent No. 6514948
 ; GENERAL INFORMATION:
 ; APPLICANT: Raz, Eyal R.
 ; TITLE OF INVENTION: METHOD FOR ENHANCING AN IMMUNE RESPONSE
 ; FILE REFERENCE: 30448-64US01
 ; CURRENT APPLICATION NUMBER: US/09/347,343A
 ; CURRENT FILING DATE: 1999-07-02
 ; NUMBER OF SEQ ID NOS: 40
 ; SOFTWARE: FastSEQ for Windows Version 3.0
 SEQ ID NO 32
 LENGTH: 22
 TYPE: DNA
 ORGANISM: synthetic oligonucleotide

Query Match 85.5%; Score 18.8; DB 4; Length 22;
 Best Local Similarity 90.5%; Pred. No. 6.3;
 Matches 20; Conservative 0; Mismatches 2; Indels 0;
 Gaps 0;
 Db 1 TGACGTGAACTTATAGATGA 22
 1 ||||| ||||| ||||| |||||

RESULT 9
 US-09-347-343-33
 ; Sequence 33, Application US/09347343A
 ; Patent No. 6534062
 ; GENERAL INFORMATION:
 ; APPLICANT: Raz, Eyal

Query Match 85.5%; Score 18.8; DB 4; Length 22;
 Best Local Similarity 90.9%; Pred. No. 6.3;
 Matches 20; Conservative 0; Mismatches 2; Indels 0;
 Gaps 0;
 Db 1 TGACGTGAACTTATAGATGA 22
 1 ||||| ||||| ||||| |||||
 RESULT 11
 US-09-820-484-3
 ; Sequence 3, Application US/09820484
 ; Patent No. 6534062
 ; GENERAL INFORMATION:
 ; APPLICANT: Raz, Eyal

Query Match 85.5%; Score 18.8; DB 4; Length 22;
 Best Local Similarity 90.9%; Pred. No. 6.3;
 Matches 20; Conservative 0; Mismatches 2; Indels 0;
 Gaps 0;
 Organism: synthetic oligonucleotide
 US-09-347-343-33
 SEQ ID NO 33
 LENGTH: 22
 TYPE: DNA
 ORGANISM: synthetic oligonucleotide
 US-09-820-484-1
 ; Sequence 1, Application US/09820484
 ; Patent No. 6534062
 ; GENERAL INFORMATION:
 ; APPLICANT: Raz, Eyal
 ; TITLE OF INVENTION: Method for Increasing a Cytotoxic T
 ; TITLE OF INVENTION: Lymphocyte Response in vivo.
 ; FILE REFERENCE: 06510-188US1
 ; CURRENT APPLICATION NUMBER: US/09/820,484
 ; CURRENT FILING DATE: 2000-03-18
 ; PRIOR APPLICATION NUMBER: US 60/192,537
 ; PRIOR FILING DATE: 2000-01-28
 ; PRIOR APPLICATION NUMBER: US 60/203,567
 ; PRIOR FILING DATE: 2000-05-11
 ; PRIOR APPLICATION NUMBER: US 60/215,895
 ; PRIOR FILING DATE: 2000-07-05
 ; NUMBER OF SEQ ID NOS: 8
 ; SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 1
 LENGTH: 22
 TYPE: DNA
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: Disulfide-linked phosphothioate ISS-ODN
 NAME/KEY: modified base
 LOCATION: (1) ... (1)
 OTHER INFORMATION: disulfide thymine
 US-09-820-484-1
 ; Sequence 3, Application US/09820484
 ; Patent No. 6534062
 ; GENERAL INFORMATION:
 ; APPLICANT: Raz, Eyal

APPLICANT: Cho, Hearn Jay
 APPLICANT: Richman, Anthony A.
 APPLICANT: Horner, Douglas
 TITLE OF INVENTION: Method for Increasing a Cytotoxic T Lymphocyte Response in vivo.
 FILE REFERENCE: 06510-188US1
 CURRENT APPLICATION NUMBER: US/09/820,484
 CURRENT FILING DATE: 2001-03-28
 PRIOR APPLICATION NUMBER: US 60/192,537
 PRIOR FILING DATE: 2000-03-28
 PRIOR APPLICATION NUMBER: US 60/203,567
 PRIOR FILING DATE: 2000-05-11
 PRIOR APPLICATION NUMBER: US 60/215,895
 PRIOR FILING DATE: 2000-07-05
 NUMBER OF SEQ ID NOS: 8
 SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 3
 LENGTH: 22
 TYPE: DNA
 ORGANISM: Artificial Sequence
 FEATURE: phosphorothioate ISS-ODN
 OTHER INFORMATION: US-09-820-484-3

Query Match Score 18.8; DB 4; Length 22;
 Best Local Similarity 90.9%; Pred. No. 6.3;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 Qy 1 TGACTGTGAACTTATAGATGA 22
 Db 1 TGACTGTGAACTTCGGATGA 22

RESULT 12
 US-09-820-484-7
 ; Sequence 7, Application US/09820484
 ; Patent No. 6534062
 ; GENERAL INFORMATION:
 ; APPLICANT: Raz, Byal
 ; APPLICANT: Cho, Hearn Jay
 ; APPLICANT: Richman, Anthony A.
 ; APPLICANT: Horner, Douglas
 TITLE OF INVENTION: Method for Increasing a Cytotoxic T Lymphocyte Response in vivo.
 FILE REFERENCE: 06510-188US1
 CURRENT APPLICATION NUMBER: US/09/820,484
 CURRENT FILING DATE: 2001-03-28
 PRIOR APPLICATION NUMBER: US 60/192,537
 PRIOR FILING DATE: 2000-03-28
 PRIOR APPLICATION NUMBER: US 60/203,567
 PRIOR FILING DATE: 2000-05-11
 PRIOR APPLICATION NUMBER: US 60/215,895
 PRIOR FILING DATE: 2000-07-05
 NUMBER OF SEQ ID NOS: 8
 SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 7
 LENGTH: 22
 TYPE: DNA
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: mODN
 US-09-820-484-7

Query Match Score 18.8; DB 4; Length 22;
 Best Local Similarity 90.9%; Pred. No. 6.3;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 Qy 1 TGACTGTGAACTTATAGATGA 22
 Db 1 TGACTGTGAACTTCGGATGA 22

RESULT 14
 US-09-774-403A-3
 ; Sequence 3, Application US/09774403A
 ; Patent No. 6552006
 ; GENERAL INFORMATION:
 ; APPLICANT: Raz, Byal
 ; APPLICANT: Richard Kornbluth
 ; APPLICANT: Antonio Catanzaro
 ; APPLICANT: Tomoko Hayashi
 ; APPLICANT: Dennis Carson
 TITLE OF INVENTION: Immunomodulatory Polynucleotides in Treatment of Infection by an Intracellular Pathogen
 FILE REFERENCE: UCAL166
 CURRENT APPLICATION NUMBER: US/09/774,403A
 CURRENT FILING DATE: 2002-04-15
 PRIOR APPLICATION NUMBER: 60/179,353
 PRIOR FILING DATE: 2000-01-31
 NUMBER OF SEQ ID NOS: 7
 SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 1
 LENGTH: 22
 TYPE: DNA
 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: Immunomodulatory sequence
 US-09-774-403A-1

Query Match Score 18.8; DB 4; Length 22;
 Best Local Similarity 90.9%; Pred. No. 6.3;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 Qy 1 TGACTGTGAACTTATAGATGA 22
 Db 1 TGACTGTGAACTTCGGATGA 22

RESULT 15
 US-09-774-403A-3
 ; Sequence 3, Application US/09774403A
 ; Patent No. 6552006
 ; GENERAL INFORMATION:
 ; APPLICANT: Raz, Byal
 ; APPLICANT: Richard Kornbluth
 ; APPLICANT: Antonio Catanzaro
 ; APPLICANT: Tomoko Hayashi
 ; APPLICANT: Dennis Carson
 TITLE OF INVENTION: Immunomodulatory Polynucleotides in Treatment of Infection by an Intracellular Pathogen
 FILE REFERENCE: UCAL166
 CURRENT APPLICATION NUMBER: US/09/774,403A
 CURRENT FILING DATE: 2002-04-15
 PRIOR APPLICATION NUMBER: 60/179,353
 PRIOR FILING DATE: 2000-01-31
 NUMBER OF SEQ ID NOS: 7
 SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 3
 LENGTH: 22
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 ORGANISM: Artificial Sequence
 FEATURE:
 OTHER INFORMATION: Control sequence
 US-09-774-403A-3

Query Match Score 18.8; DB 4; Length 22;
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 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 Qy 1 TGACTGTGAACTTATAGATGA 22
 Db 1 TGACTGTGAACTTCGGATGA 22

Sequence 2, Application US/09296477A
Patent No. 6589940
GENERAL INFORMATION:
APPLICANT: RAZ, E.
APPLICANT: SCHWARTZ, D.
APPLICANT: ROMAN, M.
APPLICANT: DINA, D.
TITLE OF INVENTION: IMMUNOSTIMULATORY OLIGONUCLEOTIDES, COMPOSITIONS THEREOF AND METHODS OF USE
TITLE OF INVENTION: THEREOF
FILE REFERENCE: 37788200420
CURRENT APPLICATION NUMBER: US/09/296,477A
CURRENT FILING DATE: 1999-04-22
EARLIER APPLICATION NUMBER: 09/092,329
EARLIER FILING DATE: 1998-06-05
EARLIER APPLICATION NUMBER: 60/048,793
EARLIER FILING DATE: 1997-06-06
NUMBER OF SEQ ID NOS: 21
SOFTWARE: FastSEQ for Windows Version 3.0
SEQ ID NO 2
LENGTH: 22
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic construct
US-09-296-477-2

Query Match 85 5%; Score 18.8; DB 4; Length 22;
Best Local Similarity 90.9%; Pred. No. 6.3;
Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Db 1 TGACTGTGAACGTTCGAGATCA 22

Search completed: November 9, 2005, 19:20:07
Job time : 96 secs

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					SOURCE	synthetic construct	
					ORGANISM	other sequences; artificial sequences.	
					REFERENCE	1 (bases 1 to 22)	
					AUTHORS	Carpentier,A.	
					TITLE	Use of stabilized oligonucleotide for producing agents having antitumor activity	
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					PF	17-MAR-2000 JP 20000606246	
					PR	19-MAR-1999 FR 99/03433	
					PI	ANTOINE CARBENTIER	
					PC	AS1K47/48, A61K31/711, A61P35/00	
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DEFINITION	AX036952.		AUTHORS Raz, E. and Roman, M.
ACCESSION	AX036952.		TITLE Patent: JP 200505580-A 2 19-FEB-2002; DYNAVAX TECHNOLOGIES CORP., THE REGENTS OF THE UNIVERSITY OF CALIFORNIA
VERSION	1	GI:11226380	JOURNAL OS Artificial Sequence
KEYWORDS			PN JP 200250580-A/2
SOURCE			PD 19-FEB-2002
ORGANISM			PF 05-JUN-1998 JP 1999502803
REFERENCE			PR 06-JUN-1997 US 60/048793
AUTHORS			PI EYAL RAZ, MARK ROMAN
JOURNAL			PC C12N15/00, C12N15/63, C12N15/79, C12N15/09, A61K48/00 CC Oligonucleotide
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Db	1	TGACTGTGAACTTATAGATGA 22	Db 1 TGACTGTGAACTTATAGATGA 22
RESULT 3	AR148608	AR148608 Sequence 2 from patent US 6225292.	RESULT 5
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VERSION	AR148608.1	linear	LOCUS Sequence 5 from patent US 6613751.
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SOURCE	Unknown.		SEQUENCE
ORGANISM	Unclassified.		ACCESSION AR392166
	1 (bases 1 to 22)		VERSION AR392166.1 GI:40116143
REFERENCE	Raz, E. and Roman, M.		KEYWORDS Unknown.
AUTHORS	Inhibitors of DNA immunostimulatory sequence activity		SOURCE Unknown.
TITLE	Patent: US 6225292-A 2 01-MAY-2001; Location/Qualifiers		ORGANISM Unclassified.
JOURNAL	1..22		REFERENCE 1 (bases 1 to 22)
FEATURES	/organism="unknown"		AUTHORS Raz, E. and Rachmilewitz, D.
source	/mol_type="unassigned DNA"		TITLE Method for treating inflammatory bowel disease and other forms of
			JOURNAL Gastrointestinal inflammation
			FEATURES Patent: US 6613751-A 5 02-SEP-2003; Location/Qualifiers
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Db	1	TGACTGTGAACTTATAGATGA 22	Db 1 TGACTGTGAACTTATAGATGA 22
RESULT 4	BD13175	BD13175 Inhibitors of DNA immunostimulatory sequence activity	RESULT 6
DEFINITION	BD13175	DNA	AR392167 AR392167
ACCESSION	BD13175	linear	LOCUS Sequence 6 from patent US 6613751.
VERSION	BD13175.1		DEFINITION 22 bp
KEYWORDS	GI:23231120		SEQUENCE
SOURCE	JP 200250580-A/2.		ACCESSION AR392167
ORGANISM	Synthetic construct		VERSION AR392167.1 GI:40116144
	Other sequences: artificial sequences		KEYWORDS Unknown.
			SOURCE Unknown.
			ORGANISM Unclassified.

REFERENCE	1 (bases 1 to 22)	Qy	1 TGACTGTGAACGTTATAGATCA 22
AUTHORS	Raz, E. and Rachmowitz, D.	Db	1 TGACTGTGAACGTTATAGATCA 22
TITLE	Method for treating inflammatory bowel disease and other forms of		
JOURNAL	gastrointestinal inflammation		
FEATURES	Patent: US 6613751-A 02-SEP-2003;		
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Db	1 TGACTGTGAACGTTAGAGATCA 22		
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DEFINITION	Query Match	85.5%; Score 18.8; DB 6; Length 22;	
ACCESSION	AR148607		
VERSION	GI:15112697		
KEYWORDS			
SOURCE	Unknown.		
ORGANISM	Unclassified.		
REFERENCE	1 (bases 1 to 22)		
AUTHORS	Raz, E. and Roman, M.		
TITLE	Inhibitors of DNA immunostimulatory sequence activity		
JOURNAL	Patent: US 6225292-A 01-MAY-2001;		
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Qy	1 TGACTGTGAACGTTATAGATCA 22		
Db	1 TGACTGTGAACGTTAGAGATCA 22		
RESULT 8			
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DEFINITION	Query Match	85.5%; Score 18.8; DB 6; Length 22;	
ACCESSION	AR148609		
VERSION	GI:15112699		
KEYWORDS			
SOURCE	Unknown.		
ORGANISM	Unclassified.		
REFERENCE	1 (bases 1 to 22)		
AUTHORS	Raz, E. and Roman, M.		
TITLE	Inhibitors of DNA immunostimulatory sequence activity		
JOURNAL	Patent: US 6225292-A 01-MAY-2001;		
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Query Match	85.5%; Score 18.8; DB 6; Length 22;		
Best Local Similarity	90.9%; Pred. No. 55;		
Matches	20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;		
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SOURCE	synthetic construct	Matches	20; Conservative 0; Mismatches	
ORGANISM	other sequences; artificial sequences.	Qy	1 TGACTGTAACGTTAGATGA 22	
REFERENCE	Raz, E. and Roman, M.	Db	1 TGACTGTAACGTTAGATGA 22	
AUTHORS	Inhibitors of DNA immunostimulatory sequence activity	COMMENT		
TITLE	Patent: JP 2002505580-A 3 19-FEB-2002;	OS	Artificial Sequence	RESULT 13
JOURNAL	DYNAVAX TECHNOLOGIES CORP., THE REGENTS OF THE UNIVERSITY OF CALIFORNIA	PN	JP 2002505580-A/3	BD182369
COMMENT		PD	19-FEB-2002	Anti-tumor antigens or their epitopes against HTLV-1 tumor.
		PP	05-JUN-1998	Query Match
		PR	06-JUN-1997	Best Local Similarity
		PI	EYAL RAZ, MARK ROMAN	85.5%; Score 18.8; DB 6; Length 22;
		PC	C12N15/00, C12N15/63, C12N15/09, A61K48/00 CC	Best Local Similarity
		Oligonucleotide		90.9%; Pred. No. 55;
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ORIGIN		PP	02-NOV-2002	Query Match
		PR	08-MAY-2001	Best Local Similarity
		PI	SHINO HANABUCHI, TAKASHI OHASHI, MARI KANNAGI	85.5%; Score 18.8; DB 6; Length 22;
		PC	GOIN33/50, GOIN33/15, A61K39/00	Best Local Similarity
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		PD	14-NOV-2002	Anti-tumor antigens or their epitopes against HTLV-1 tumor.
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LOCUS	BD136183.1 GI:23231128	Matches	20; Conservative 0; Mismatches	VERSION
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VERSION	BD136183.1 GI:23231128	Db	1 TGACTGTAACGTTAGATGA 22	synthetic construct
KEYWORDS	JP 2002505580-A/10.	COMMENT		ORGANISM
SOURCE	synthetic construct	OS	Artificial Sequence	other sequences; artificial sequences.
ORGANISM	other sequences; artificial sequences.	PN	JP 2002505580-A/10	REFERENCE
REFERENCE	Raz, E. and Roman, M.	PD	19-FEB-2002	1 (bases 1 to 22)
AUTHORS	Inhibitors of DNA immunostimulatory sequence activity	PP	05-JUN-1998	Anti-tumor antigens or their epitopes against HTLV-1 tumor.
TITLE	Patent: JP 2002505580-A 10 19-FEB-2002;	PI	EYAL RAZ, MARK ROMAN	Authors: Hanabuchi, S., Ohashi, T. and Kannagi, M.
JOURNAL	DYNAVAX TECHNOLOGIES CORP., THE REGENTS OF THE UNIVERSITY OF CALIFORNIA	PC	C12N15/00, C12N15/63, C12N15/09, A61K48/00 CC	Title: JPN SCIENCE AND TECHNOLOGY CORP, SHINO HANABUCHI, TAKASHI OHASHI, MARI KANNAGI
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AUTHORS	Inhibitors of DNA immunostimulatory sequence activity	PP	05-JUN-1998	
TITLE	Patent: JP 2002505580-A 1 14-NOV-2002;	PI	SHINO HANABUCHI, TAKASHI OHASHI, MARI KANNAGI	
JOURNAL	DYNAVAX TECHNOLOGIES CORP, SHINO HANABUCHI, TAKASHI OHASHI, MARI KANNAGI	PC	GOIN33/50, GOIN33/15, A61K39/00	
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REFERENCE	Raz, E. and Roman, M.	PD	14-NOV-2002	
AUTHORS	Inhibitors of DNA immunostimulatory sequence activity	PP	05-JUN-1998	
TITLE	Patent: JP 2002505580-A 1 14-NOV-2002;	PI	EYAL RAZ, MARK ROMAN	
JOURNAL	DYNAVAX TECHNOLOGIES CORP, SHINO HANABUCHI, TAKASHI OHASHI, MARI KANNAGI	PC	GOIN33/50, GOIN33/15, A61K39/00	
COMMENT		FH	Key	
		FT	source	

JOURNAL Patent: JP 2002372532-A 1 26-DEC-2002;
 COMMENT OS JAPAN SCIENCE AND TECHNOLOGY CORP
 OS Artificial Sequence
 PN JP 2002372532-A/1
 PD 26-DEC-2002
 PP 08-MAY-2001 JP 2001137526
 PI SHINO, HANABUCHI, TAKASHI, OHASHI, MARI, KANNAGI
 G01N3/3/50, A61K39/00, A61K39/21, A61P35/00, A61P35/02, A61P37/04,
 PC C07K7/06,
 PC C12N5/06, C12Q1/02, G01N33/00, G01N33/53, G01N33/53, PC
 G01N33/56,
 PC G01N33/574
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 Best Local Similarity 90.9%; Pred. No. 55;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 KEYWORDSDb 1 TGACTGTGAACTTATAGATGA 22
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 DEFINITION BD190435
 ACCESSION BD190435
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 SOURCE synthetic construct
 ORGANISM synthetic construct
 OTHER SEQUENCES; artificial sequences.
 REFERENCE 1 (bases 1 to 22)
 AUTHORS Barackman, J., Simph, M., Ugozoli, M., Kazazu, J., Donnelly, J.,
 Ott, G.S. and Ohagan, D.
 TITLE Microemulsions with Adsorbed Macromolecules and Microparticles
 JOURNAL Chiron Corporation
 COMMENT OS Artificial Sequence
 PN JP 2002537102-A/19
 PD 05-NOV-2002
 PP 09-FEB-2000 JP 20000600618
 PR 29-JUL-1999 US 60/146391, 28-OCT-1999 US 60/161997, PR
 26-FEB-1999 US 60/121858
 PI John barackman, mammohan simph, mildred ugozoli, jina kazazu, john
 donnnelly,
 PI gary s ort, derek ohagan
 CC Oligonucleotide
 FH Key Location/Qualifiers
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 Best Local Similarity 90.9%; Pred. No. 55;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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		35	18.8	85.5	22	3	Aac64051	Immunost
		36	18.8	85.5	22	4	Aah2044	Cpg motif
		37	18.8	85.5	22	4	Aah20403	Cpg motif
		38	18.8	85.5	22	4	Aah43345	Immunomod
		39	18.8	85.5	22	4	Aah43340	Immunomod
		40	18.8	85.5	22	4	Aah43338	Immunomod
		41	18.8	85.5	22	4	Aah73439	Immunomod
		42	18.8	85.5	22	4	Aah73441	Immunomod
		43	18.8	85.5	22	4	Aah75992	Immunomod
		44	18.8	85.5	22	4	Aai164301	Control O
		45	18.8	85.5	22	4	Aah75995	Immunomod
							Aah76000	Control O

Maximum Match 100%

```

Maximum Match 100%
Listing first 45 summaries

database : N_Geneseq1.6Dec04:*
 1: geneseqn1908s:*
 2: geneseqn1908s:*
 3: Geneseqn2000s:*
 4: Geneseqn2001as:*
 5: geneseqn2001bs:*
 6: Geneseqn2002as:*
 7: geneseqn2002bs:*
 8: geneseqn2003as:*
 9: Geneseqn2003bs:*
10: Geneseqn2003cb:*
11: Geneseqn2003ds:*
12: Geneseqn2004as:*
13: Geneseqn2004bs:*

```

uclonucleotide with antitumour activity.

古漢山脈之植物

* * *

sec04: *
sec08: *
sec908: *
sec008: *
sec01as: *
sec01bs: *
sec02as: *
sec02bs: *
sec03as: *
sec03bs: *
sec003CB: *
sec003db: *
sec00435: *

Genes genes genes genes genes genes genes genes genes gene gene gene

N_C
1: 2: 3: 4: 5: 6: 7: 8: 9: 10: 11: 12:

abage

at the

SUMMARIES						
Result No.	Score	Query Match	Length	DB ID	Description	
1	22	100.0	22	3	AAA36260 Sequence	PD 28-SEP-2000.
2	20.4	92.7	22	4	AHH43343 Immunomod	XX XX 17-MAR-2000; 2000WO-FR000676.
3	20.4	92.7	22	4	AHH43342 Immunomod	XX PR 19-MAR-1999; 99FR-0003433.
4	20.4	92.7	22	6	AAD24895 Methylate	XX XX (ASSI-) ASSISTANCE PUBLIQUE HOPITAUX PARIS.
5	20.4	92.7	22	6	AAD24894 Immunosti	XX (INRM) INST NAT SANTE & RECH MEDICALE.
6	20.4	92.7	22	12	AD055351 Immune mo	XX Carpenter A;
7	19.4	88.2	22	12	AD055287 Immune mo	XX DR WPI; 2000-602192/57.
8	18.8	85.5	22	2	AAY32079 Nucleotid	XX Use of stabilized oligonucleotides as antitumor agents, particularly
9	18.8	85.5	22	2	Aav80099 Immunomod	XX against nervous system tumors, have optimal activity and are not toxic.
10	18.8	85.5	22	2	Aav80097 Immunomod	XX
11	18.8	85.5	22	2	Aav80103 Immunomod	XX Claim 3; Page 48; 57pp; French.
12	18.8	85.5	22	2	Aav80106 Oligo use	XX
13	18.8	85.5	22	2	Aav80101 Immunomod	XX The present sequence represents a stabilised oligonucleotide which has
14	18.8	85.5	22	2	Aav80104 Oligo use	XX antitumour activity. The oligonucleotide comprises an octomer motif of
15	18.8	85.5	22	2	Aav80102 Immunomod	XX the type 5'-purine-purine-C-pyrimidine-X-X-3', where the pair
16	18.8	85.5	22	2	Aav55797 Immunosti	XX X-X is AT, AA, CT or TT. The oligonucleotides are immunomodulatory, and
17	18.8	85.5	22	2	Aav55788 Immunosti	XX are not toxic. They may be adapted for use in animals or humans. The
18	18.8	85.5	22	2	Aav55790 Immunosti	XX stabilised oligonucleotides are used for treating tumours, of any type
19	18.8	85.5	22	2	Aax336624 ISS-ODN D	XX and any degree of anaplasia, particularly glioblastomas, medulloblastomas,
20	18.8	85.5	22	3	AAA14469 Mutant im	XX or central nervous systems.

CC neuroblastomas, melanomas or carcinomas
 XX Sequence 22 BP; 7 A; 2 C; 6 G; 7 T; 0 U; 0 Other;
 SQ Query Match 100.0%; Score 22; DB 3; Length 22;
 Best Local Similarity 100.0%; Pred. No. 0.65; Indels 0; Gaps 0;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 XX

RESULT 2
 AAH43343 standard; DNA; 22 BP.
 XX
 ID AAH43343
 AC AAH43343;
 XX
 DT 13-DEC-2001 (first entry)
 XX
 DE Immunomodulatory polynucleotide 1039.
 XX
 KW Immunomodulation; Gastrointestinal tract;
 KW ulcerative colitis; Crohn's disease; inflammatory bowel disease;
 KW diarrhoea; rectal bleeding; weight loss; colon; lesion; ss.
 XX
 OS Synthetic.
 XX
 PN WO200162207-A2.
 XX
 PD 30-AUG-2001.
 XX
 DT 22-FEB-2001; 2001WO-US006034.
 XX
 PR 23-FEB-2000; 2000US-0184256P.
 XX
 PA (REGC) UNIV CALIFORNIA.
 XX
 PT Ameliorating gastrointestinal inflammation e.g. inflammatory bowel
 disease involves administering an immunomodulatory nucleic acid.
 XX
 PI Raz E, Rachmiliwitz D;
 XX
 DR WPI; 2001-565393/63.
 XX
 PT Ameliorating gastrointestinal inflammation e.g. inflammatory bowel
 disease involves administering an immunomodulatory nucleic acid.
 XX
 PS Claim 7; Page 28; 58pp; English.
 XX
 PT The sequences given in AAH43338-48 represent immunomodulatory
 CC polynucleotides which may be used to ameliorate inflammation of the
 CC gastrointestinal tract by administering a nucleic acid comprising one of
 CC these sequences. These polynucleotides all comprise an immunomodulatory
 CC nucleotide sequence of 5'-CpG-3', (I). The nucleotides may be used for
 CC ameliorating or reducing gastrointestinal inflammation e.g. chronic or
 CC acute gastrointestinal inflammation, ulcerative colitis, Crohn's disease
 CC caused by inflammatory bowel disease; diarrhoea, rectal bleeding, weight
 CC loss; to reduce colon weight and colon lesions; to reduce a colonic
 CC inflammation. The immunomodulatory polynucleotides treat inflammatory
 CC bowel disease satisfactorily and effectively and have little or no
 CC toxicity even at a high dosage of 50000 micro-g. They also reduce the
 CC risk of colonic cancer by treating ulcerative colitis
 XX
 SQ Sequence 22 BP; 7 A; 2 C; 7 G; 6 T; 0 U; 0 Other;
 CC Query Match 92.7%; Score 20.4; DB 4; Length 22;
 CC Best Local Similarity 95.5%; Pred. No. 3.9;
 CC Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 CC DE Methylated (5-methyl C) immunostimulatory oligodeoxynucleotide (ISS-ODN).
 CC XX Cell death; DNA damage; DNA-dependent protein kinase; DNA-PK; necrosis;
 CC immune response; apoptosis; Alzheimer's disease; Parkinson's disease;
 CC rheumatoid arthritis; inflammation; osteoporosis; myocardial infarction;
 CC KW

RESULT 3
 AAH43342 standard; DNA; 22 BP.
 ID AAH43342
 XX
 AC AAH43342;
 XX
 DT 13-DEC-2001 (first entry)
 XX
 DE Immunomodulatory polynucleotide 1038.
 XX
 KW Immunomodulation; inflammation; gastrointestinal tract;
 KW ulcerative colitis; Crohn's disease; inflammatory bowel disease;
 KW diarrhoea; rectal bleeding; weight loss; colon; lesion; ss.
 XX
 OS Synthetic.
 XX
 PN WO200162207-A2.
 XX
 PD 30-AUG-2001.
 XX
 DT 22-FEB-2001; 2001WO-US006034.
 XX
 PR 23-FEB-2000; 2000US-0184256P.
 XX
 PA (REGC) UNIV CALIFORNIA.
 XX
 PT Ameliorating gastrointestinal inflammation e.g. inflammatory bowel
 disease involves administering an immunomodulatory nucleic acid.
 XX
 PI Raz E, Rachmiliwitz D;
 XX
 DR WPI; 2001-565393/63.
 XX
 PT Ameliorating gastrointestinal inflammation e.g. inflammatory bowel
 disease involves administering an immunomodulatory nucleic acid.
 XX
 PS Example 2; Page 28; 58pp; English.
 XX
 PT The sequences given in AAH43338-48 represent immunomodulatory
 CC polynucleotides which may be used to ameliorate inflammation of the
 CC gastrointestinal tract by administering a nucleic acid comprising one of
 CC these sequences. These polynucleotides all comprise an immunomodulatory
 CC nucleotide sequence of 5'-CpG-3', (I). The nucleotides may be used for
 CC ameliorating or reducing gastrointestinal inflammation e.g. chronic or
 CC acute gastrointestinal inflammation, ulcerative colitis, Crohn's disease
 CC caused by inflammatory bowel disease; diarrhoea, rectal bleeding, weight
 CC loss; to reduce colon weight and colon lesions; to reduce a colonic
 CC inflammation. The immunomodulatory polynucleotides treat inflammatory
 CC bowel disease satisfactorily and effectively and have little or no
 CC toxicity even at a high dosage of 50000 micro-g. They also reduce the
 CC risk of colonic cancer by treating ulcerative colitis
 XX
 SQ Sequence 22 BP; 7 A; 2 C; 7 G; 6 T; 0 U; 0 Other;
 CC Query Match 92.7%; Score 20.4; DB 4; Length 22;
 CC Best Local Similarity 95.5%; Pred. No. 3.9;
 CC Matches 21; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 CC DE Methylated (5-methyl C) immunostimulatory oligodeoxynucleotide (ISS-ODN).
 CC XX Cell death; DNA damage; DNA-dependent protein kinase; DNA-PK; necrosis;
 CC immune response; apoptosis; Alzheimer's disease; Parkinson's disease;
 CC rheumatoid arthritis; inflammation; osteoporosis; myocardial infarction;
 CC KW

RESULT 4
 AAD24895 standard; DNA; 22 BP.
 ID AAD24895
 XX
 AC AAD24895;
 XX
 DT 12-MAR-2002 (first entry)
 XX
 DE Methylated (5-methyl C) immunostimulatory oligodeoxynucleotide (ISS-ODN).
 XX
 KW Cell death; DNA damage; DNA-dependent protein kinase; DNA-PK; necrosis;
 KW immune response; apoptosis; Alzheimer's disease; Parkinson's disease;
 KW rheumatoid arthritis; inflammation; osteoporosis; myocardial infarction;

Query 1 TGACTGTGAACTTATAGATGA 22
 DB 1 TGACTGTGAACTTATAGATGA 22

liver disease; reperfusion injury; carcinoma; multiple sclerosis; stroke; amyotrophic lateral sclerosis; Acquired Immune Deficiency Syndrome; AIDS; head injury damage; aplastic anaemia; tumour; organ transplantation; cerebral infarction; follicular lymphomas; systemic lupus erythematosus; viral infection; Glomerulonephritis; apoptosis; apoptosis; autoimmune disorder; sepsis; immunostimulatory oligodeoxynucleotide; ISS-ODN; ss.	
Unidentified.	
OS	Location/Qualifiers
XX	
KEY	11
modified_base	
FT	/*tag= a
FT	/mod_base= m5C
XX	
PN	
PN	
XX	
XX	WO200185910-A2.
XX	
PD	15-NOV-2001.
XX	
XX	04-MAY-2001; 2001WO-US14508.
XX	
PR	05-MAY-2000; 2000US-0202274P.
PR	
XX	17-JAN-2001; 2001US-0262321P.
PA	
PA	(REGC) UNIV CALIFORNIA.
XX	
XX	Raz E, Lois AF, Takabayashi K;
XX	
WPTI	2002-062244/08.
XX	
PPT	Modulating cell death or reducing DNA damage in eukaryotic cells, useful for reducing cell death in individual or organ, comprises contacting cell with agent modulating biological activity of DNA-dependent protein
PPT	

Example 3: Page 33; 57pp; English.

THE invention relates to a method for modulating cell death or reducing DNA damage in an eukaryotic cell by contacting the cell with an agent that modulates the biological activity of DNA-dependent protein kinase (DNA-PK). The invention also relates nucleic acids which modulate the immune response binding to KU antigen, resulting in activation of DNA-PK. The method is useful for modulating cell death or reducing DNA damage in an eukaryotic cell, for treating any disorder resulting from a genotoxic agent inserted to a cell e.g., necrosis, apoptosis. The method is also useful for treating cell death-related indications such as Alzheimer's disease, Parkinson's disease, rheumatoid arthritis, septic shock, sepsis, stroke, central nervous system inflammation, osteoporosis, degenerative liver disease, cerebellar degeneration, reperfusion injury, multiple sclerosis, lateral sclerosis, myocardial infarction, head injury, damage, acquired immunodeficiency syndrome (AIDS), aplastic anaemia, cerebral infarction, bypass heart surgery, organ transplantation. The method is also useful for treating follicular lymphomas, carcinomas, autoimmune disorders (systemic lupus erythematosus), hormone dependent tumours, immunemediated glomerulonephritis; apoptosis and viral infections. The present sequence is methylated (5-methyl C) immunostimulatory oligodeoxynucleotide (ISS-ODN) used in the exemplification of the invention.

Sequence 22 BP; 7 A; 2 C; 7 G; 6 T; 0 U; 0 Other;
Sequence 22 BP; 7 A; 2 C; 7 G; 6 T; 0 U; 0 Other;

Qy	1	TGACCTGTGAACGTTATAGATCA	22
pb	1	TGACCTGTGAACGTTATAGATCA	22

Best Local Similarity 95.5%; Fred. No. 3.9;
Matches 21; Conservative 0; Mismatches 1; Indels

```

Query Match      92.7%;  Score 20.4;  DB 6;  Length 22;
Best Local Similarity 95.5%;  Pred. No. 3.9;
Matches 21;  Conservative 0;  Mismatches 1;  Indels 0;  Gaps 0;
Qy  1 TGACTCTGAAACGTTATAGATGAA 22

```

AC	AA24894;
XX	12-MAR-2002 (first entry)
XX	Immunostimulatory oligodeoxynucleotide (ISS-ODN) 2.
XX	Cell death; DNA damage; DNA-dependent protein kinase; DNA-PK; necrosis; immune response; apoptosis; Alzheimer's disease; Parkinson's disease; rheumatoid arthritis; inflammation; osteoporosis; myocardial infarction; liver disease; reperfusion injury; carcinoma; multiple sclerosis; stroke; amyotrophic lateral sclerosis; Acquired Immune Deficiency Syndrome; AIDS; head injury damage; aplastic anemia; tumor; organ transplantation; cerebral infarction; follicular lymphomas; systemic lupus erythematosus; viral infection; glomerulonephritis; apoptosis; autoimmunity disorder; sepsis; immunostimulatory oligodeoxynucleotide; ISS-ODN; 88.
DB	
DE	
KW	

Unidentified.
WO2000185910-A2.
115.NOV-2001.
004-MAY-2001; 2001WO-US014508.
005-MAY-2000; 2000US-020274P.
117.JAN-2001; 2001US-0262321P.
(RGSC) UNIV CALIFORNIA.
Raz E, Lois AF, Takabayashi K;
WPP; 2002-052244/08.
Modulating cell death or reducing DNA damage in eukaryotic cells, useful
for reducing cell death in individual or organ, comprises contacting cell
with agent modulating biological activity of DNA-dependent protein

Example 3: Page 33; 57pp; English.

The invention relates to a method for modulating cell death or reducing DNA damage in an eukaryotic cell by contacting the cell with an agent that modulates the biological activity of DNA-dependent protein kinase (DNA-PK). The invention also relates nucleic acids which modulate the immune response binding to Ku antigen, resulting in activation of DNA-PK. The method is useful for modulating cell death or reducing DNA damage in an eukaryotic cell, for treating any disorder resulting from a genotoxic insult to a cell e.g., necrosis, apoptosis. The method is also useful for treating cell death-related indications such as Alzheimer's disease, Parkinson's disease, rheumatoid arthritis, septic shock, stroke, central nervous system inflammation, osteoporosis, degenerative liver disease, cerebellar degeneration, reperfusion injury, multiple sclerosis, reperfusion injury, myocardial infarction, head injury, damage, acquired immunodeficiency syndrome (AIDS), aplastic anaemia, cerebral infarction, bypass heart surgery, organ transplantation. The method is also useful for treating follicular lymphomas, carcinomas, autoimmune disorders (systemic lupus erythematosus), hormone dependent tumours, immunomodulatory mediated glomerulonephritis; apoptosis and viral infections. The present sequence is immunosimulatory oligodeoxynucleotide (ISS-ODN) used for identifying ISS-binding protein, which is used in the exemplification of the invention.

RESULT 6	
AD055351	AD055351. standard: DNA; 22 BP.
26-AUG-2004 (first entry)	Immune modulatory nucleic acid (IMs) #126.
Immune modulatory nucleic acid (IMs) #126.	Immune modulatory nucleic acid; IMs; immune modulatory sequence; non CpG; self-molecule related disease; autoimmune disease; multiple sclerosis; rheumatoid arthritis; insulin-dependent diabetes mellitus; Sjogren's syndrome; primary biliary cirrhosis; myasthenia gravis; autoimmune uveitis; pemphigus vulgaris; systemic lupus erythematosus; ankylosing spondylitis; autoimmune skin disease; Grave's disease; inflammatory disease; osteoarthritis; gout; pseudogout; hydroxyapatite deposition disease; asthma; bursitis; tendonitis; conjunctivitis; urethritis; cystitis; balanitis; dermatitis; spinal cord injury; peptic ulcer; hyperlipidaemia; coronary artery disease; migraine; neuroprotective; antirheumatic; antiarthritic; antidiabetic; osteopathic; antigout; antiasthmatic; antiinflammatory; ophthalmological; dermatological; vasotropic; antimigraine; vaccine; gene therapy; ss.
OS	Synthetic.
FFH	Key
FT	misc_feature
FT	feature
XX	Location/Qualifiers
XX	9..14
XX	/*tag= a
XX	/note= "Core Pu-Pu-X-Y-Py-Py hexamer region"
XX	11..12
XX	/*tag= b
XX	/note= "GpG or non-GpG, non-CpG dinucleotide"
W02004047734-A2.	
10-JUN-2004.	
21-NOV-2003.	2003WO-US037157.
21-NOV-2002.	2002US-0428643P.
(BAYH-)	BAYHILL THERAPEUTICS INC.
(STRD)	UNIV LEELAND STANFORD JUNIOR.
Garren H.	Ho PP, Steinman L;
WPI:	2004-441065/41.
XX	Pharmaceutical compositions comprising an immune modulatory nucleic acid comprising a hexamer region, useful for treating an autoimmune disease, e.g. multiple sclerosis, rheumatoid arthritis or insulin dependent diabetes mellitus.
XX	Example 10: Page 68; 90pp; English.
XX	The invention relates to a pharmaceutical composition for treating a disease associated with one or more self-molecules present non-physiologically in an individual (e.g., autoimmune diseases), comprising an immune modulatory nucleic acid (IMs, immune modulatory sequence) comprising a hexamer region of the formula 5'-purine-pyrimidine-[X]-[Y]-purine-pyrimidine-3', where X and Y are any naturally occurring or synthetic nucleotides except cytosine-guanine, and a pharmaceutical carrier. The immune modulatory nucleic acid may also contain a polyG region linked 5' and/or 3' to the hexamer region. The invention also relates to a nucleic acid composition comprising a nucleic acid vector having at least one cytosine to non-cytosine substitution (preferably C to G) within a Cpg motif, wherein the Cpg motif is of the formula: (a) 5'-purine-pyrimidine-C-G-pyrimidine-purine-pyrimidine-3'; or (b) 5'-purine-G-pyrimidine-pyrimidine-3'. The immune modulatory nucleic acid sequences are useful in the treatment of disease associated with one or more self-molecules present non-physiologically in an individual, such as those useful in the treatment of disease associated with one or more self-molecules.
OS	Synthetic.
XX	Key
XX	misc_feature
FT	feature
FT	feature
FT	feature
FT	modified_base
FT	modified_base
PN	Location/Qualifiers
XX	9..14
XX	/*tag= a
XX	/note= "Core Pu-Pu-X-Y-Py-Py hexamer region"
XX	11..12
XX	/*tag= b
XX	/note= "GpG or non-GpG, non-CpG dinucleotide"
XX	/mod_base= i
XX	WO2004047734-A2.
XX	10-JUN-2004.
XX	21-NOV-2003; 2003WO-US037157.
XX	PR 21-NOV-2002; 2002US-0428643P.
XX	PR 21-NOV-2002; 2002US-0428643P.
PA (BAYH-)	BAYHILL THERAPEUTICS INC.
PA (STRD)	UNIV LEELAND STANFORD JUNIOR.

XX	Garren H,	Ho PP,	Steinman L;	
PI				
XX				
DR	WPI: 2004-441065/41.			
XX				
PT	Pharmaceutical compositions comprising an immune modulatory nucleic acid			
PT	comprising a hexamer region, useful for treating an autoimmune disease,			
PT	e.g. multiple sclerosis, rheumatoid arthritis or insulin dependent			
PT	diabetes mellitus.			
XX				
PS	Example 10; Page 66; 98pp; English.			
XX				
CC	The invention relates to a pharmaceutical composition for treating a			
CC	disease associated with one or more self-molecules present non-			
CC	physiologically in an individual (e.g., autoimmune diseases), comprising			
CC	an immune modulatory nucleic acid (IMM, immune modulatory sequence)			
CC	comprising a hexamer region of the formula 5'-purine-pyrimidine-[X]-			
CC	pyrimidine-pyrimidine-3', where X and Y are any naturally-occurring or			
CC	synthetic nucleotides except cytosine-guanine, and a pharmaceutical			
CC	carrier. The immune modulatory nucleic acid may also contain a polyG			
CC	region linked 5' and/or 3' to the hexamer region. The invention also			
CC	relates to a nucleic acid composition comprising a nucleic acid vector			
CC	preferably C			
CC	having at least one cytosine to non-cytosine substitution (preferably C			
CC	to G) within a CpG motif, wherein the CpG motif is of the formula: (a) 5'-			
CC	purine-Pyrimidine-C-G-pyrimidine-3'; or (b) 5'-purine-purine-C			
CC	G-pyrimidine-pyrimidine-3'. The immune modulatory nucleic acid sequences			
CC	are useful in the treatment of disease associated with one or more self-			
CC	molecules present non-physiologically in an individual, such as			
CC	autoimmune diseases (e.g., multiple sclerosis, rheumatoid arthritis,			
CC	insulin-dependent diabetes mellitus, autoimmune uveitis, primary biliary			
CC	cirrhosis, myasthenia gravis, Sjogren's syndrome, pemphigus vulgaris,			
CC	scleroderma, pernicious anaemia, systemic lupus erythematosus, ankylosing			
CC	spondylitis, autoimmune skin diseases and Grave's disease); inflammatory			
CC	diseases (e.g., osteoarthritis, gout, pseudogout, hydroxyapatite			
CC	deposition disease, asthma, bursitis, tendonitis, conjunctivitis,			
CC	urethritis, cystitis, balanitis and dermatitis); or other conditions such			
CC	as spinal cord injury, peptic ulcer, hyperlipidaemia, coronary artery			
CC	disease and migraine. The present sequence represents a specific example			
CC	of an immune modulatory nucleic acid predicted to be useful for			
CC	modulating autoimmune disease which is referred to in an example of the			
CC	invention.			
XX				
SQ	Sequence 22 BP; 7 A; 1 C; 7 G; 6 T; 0 U; 1 Other;			
Query Match	88.2%;	Score 19.4;	DB 12;	Length 22;
Best Local Similarity	90.9%;	Pred. No. 12;		
Matches	20;	Conservative	0;	Indels 0; Gaps 0;
Qy	1 TGACTGTGAACGTTATAGATGA 22			
Db	1 TGACTGTGAACGTTAGATGA 22			
RESULT 8				
AAV32079	AAV32079 standard; DNA; 22 BP.			
ID				
XX				
AC	AAV32079;			
XX				
DT	09-SEP-1998 (First entry)			
XX				
DB	Nucleotide sequence of DY1018.			
XX				
KW	DY1018; beta-gal; ISS-PN/IMM; antigen; immune response; antibody;			
KW	immunisation; anaphylaxis; IGE; retinopathies; ss.			
XX				
Synthetic.				
OS				
XX				
Key	Location/Qualifiers			
modified_base	1..22			
FT	/*tag= a phosphothioate backbone */			
FT	note=			
XX				

PA (DYN-) DYNAVAX TECHNOLOGIES CORP.
 XX
 PI Schwartz D, Roman M, Dina D;
 XX
 WPI: 1999-059898/05.

PT Immunostimulatory oligonucleotides regulate the immune system - and contain an immune-stimulating octanucleotide sequence; for treating cancer, allergic and infectious diseases.
 XX
 PS Claim 8; Page 29; 63pp; English.

CC The invention relates to immunomodulatory oligonucleotides that comprise at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS sequences are selected from the group consisting of AACGTTCC, AACGTTCG, GACGTTCC, and GACGTTCG. The immunomodulatory sequences are used to treat patients needing immune regulation, such as those suffering from cancer, an allergic disease and asthma. They are also used to prevent infectious diseases such as influenza, herpes, hepatitis B, human immunodeficiency and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and Bordetella pertussis, malaria plasmodia, Leishmania, Trypanosoma and Schistosoma. The immunomodulatory sequences are used to treat patients needing immune regulation, such as those suffering from cancer, an allergic disease and asthma. They are also used to prevent infectious diseases such as influenza, herpes, hepatitis B, human immunodeficiency and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and Bordetella pertussis, malaria plasmodia, Leishmania, Trypanosoma and Schistosoma. The immunomodulatory sequences are used to screen for human immunostimulatory activity by incubating macrophage cells and the oligonucleotide; and determining the relative amount of Th1-biased cytokines in the supernatant. Sequences AAV80096 to AAV80103 represent specific claimed examples of such immunomodulatory oligonucleotides
 XX
 SQ Sequence 22 BP; 6 A; 4 C; 6 G; 6 T; 0 U; 0 Other;

Query Match 85.5%; Score 18.8; DB 2; Length 22;
 Best Local Similarity 90.9%; Pred. No. 24;
 Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Query 1 TGACTGTGAACTGTATAGATGA 22
 Db 1 TGACTGTGAACTGTCCAGATGA 22

RESULT 11
 AAV80103
 ID AAV80103 standard; DNA; 22 BP.
 XX
 AC AAV80103;
 XX
 DT 12-MAR-1999 (first entry)
 XX
 DB Immunomodulatory oligo comprising an ISS sequence.
 XX
 KW Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;
 KW ISS: cancer; allergy; asthma; hepatitis B infection; papillomavirus;
 KW human immunodeficiency virus; influenza; herpes; M. tuberculosis; B.
 KW pertussis; malaria; plasmodia; Leishmania; Trypanosoma; Schistosoma.
 XX
 OS Synthetic.
 XX
 DT 12-MAR-1999 (first entry)
 XX
 DE Immunomodulatory oligo comprising an ISS sequence.
 XX
 KW Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;
 KW ISS: cancer; allergy; asthma; hepatitis B infection; papillomavirus;
 KW human immunodeficiency virus; influenza; herpes; M. tuberculosis;
 KW B. pertussis; malaria; plasmodia; Leishmania; Trypanosoma; Schistosoma.
 OS Synthetic.
 PN WO9855495-A2.
 XX
 PD 10-DEC-1998.
 XX
 PP 05-JUN-1998; 98WO-US011578.
 XX
 PR 06-JUN-1997; 97US-004873P.
 XX
 PA (DYN-) DYNAVAX TECHNOLOGIES CORP.
 XX
 PI Schwartz D, Roman M, Dina D;
 XX
 WPI: 1999-059898/05.

PA Immunostimulatory oligonucleotides regulate the immune system - and contain an immune-stimulating octanucleotide sequence; for treating cancer, allergic and infectious diseases.
 XX
 PR The invention relates to immunomodulatory oligonucleotides that comprise at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS sequences are selected from the group consisting of AACGTTCC, AACGTTCG,
 CC

CC GACGTTTC, and GACGTTTC. The immunomodulatory sequences are used to treat patients needing immune regulation, such as those suffering from cancer, an allergic disease and asthma. They are also used to prevent infectious diseases such as influenza, herpes, hepatitis B, human immunodeficiency and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and *Bordetella pertussis*, *malaria* plasmodia, *Leishmania*, *Trypanosoma* and *Schistosoma*. The immunomodulatory sequences are used to screen for human immunomodulatory activity by incubating macrophage cells and the oligonucleotide; and determining the relative amount of Th1-biased cytokines in the supernatant. Sequences AAV80096 to AAV80103 represent specific claimed examples of such immunomodulatory oligonucleotides

XX Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 85.5%; Score 18.8; DB 2; Length 22;

Best Local Similarity 90.9%; Pred. No. 24; Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Db 1 TGACGTGAACTTATAGATGA 22

Db 1 TGACGTGAACTTCAGATGA 22

RESULT 13

AAV80101 standard; DNA; 22 BP.

Qy 1 TGACGTGAACTTATAGATGA 22

Db 1 TGACGTGAACTTCAGATGA 22

RESULT 12

AAV80106 standard; DNA; 22 BP.

XX AAV80106;

AC 12-MAR-1999 (first entry)

DT 12-MAR-1999

DE Oligo used in experiments for stimulation of cytokine production.

XX ImmunoModulatory; immunostimulatory; octanucleotide; immune regulation;

XX human immunodeficiency virus; influenza; herpes; M. tuberculosis; SB;

XX B. pertussis; malaria; plasmodia; Leishmania; Trypanosoma; Schistosoma.

OS Synthetic.

XX PN WO9855495-A2.

XX PD 10-DEC-1998.

XX PR 05-JUN-1998;

XX PF 05-JUN-1998;

XX PR 06-JUN-1997;

XX PA 97US-0048793P.

XX PA (DYN) - DYNAVAX TECHNOLOGIES CORP.

XX PI Schwartz D, Roman M, Dina D;

XX DR WPI; 1999-059898/05.

XX PS Example 1; Page 29; 63pp; English.

XX The invention relates to immunomodulatory oligonucleotides that comprise

CC at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS

CC sequences are selected from the group consisting of AACGTTCC, AACGTTG,

CC GACGTTCC, and GACGTTCG. The immunomodulatory sequences are used to treat

CC patients needing immune regulation, such as those suffering from cancer,

CC an allergic disease and asthma. They are also used to prevent infectious

CC diseases such as influenza, herpes, hepatitis B, human immunodeficiency

CC and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and

CC *Bordetella pertussis*, *malaria* plasmodia, *Leishmania*, *Trypanosoma* and

CC *Schistosoma*. The immunomodulatory sequences are used to screen for human

CC immunomodulatory activity by incubating macrophage cells and the

CC oligonucleotide; and determining the relative amount of Th1-biased

CC cytokines in the supernatant. Sequences AAV80096 to AAV80103 represent

CC specific claimed examples of such immunomodulatory oligonucleotides

CC oligonucleotides that were tested for immunostimulatory activity. These CC were used in experiments for the stimulation of cytokine production and CC were found to lack immunostimulatory activity. The invention provides CC specific claimed examples (AAV80096-103) of immunomodulatory sequences XX

SQ Sequence 22 BP; 6 A; 3 C; 7 G; 6 T; 0 U; 0 Other;

Query Match 85.5%; Score 18.8; DB 2; Length 22;

Best Local Similarity 90.9%; Pred. No. 24;

Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TGACGTGAACTTATAGATGA 22

Db 1 TGACGTGAACTTCAGATGA 22

Query Match 85.5%; Score 18.8; DB 2; Length 22;

Best Local Similarity 90.9%; Pred. No. 24;

Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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CC and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and

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CC immunomodulatory activity by incubating macrophage cells and the

CC oligonucleotide; and determining the relative amount of Th1-biased

CC cytokines in the supernatant. Sequences AAV80104 to AAV80116 represent

CC oligonucleotides that were tested for immunostimulatory activity. These

CC were used in experiments for the stimulation of cytokine production and

CC were found to lack immunostimulatory activity. The invention provides

CC specific claimed examples (AAV80096-103) of immunomodulatory sequences

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Query Match 85.5%; Score 18.8; DB 2; Length 22;

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Matches 20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TGACGTGAACTTATAGATGA 22

Db 1 TGACGTGAACTTCAGATGA 22

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Db 1 TGACGTGAACTTCAGATGA 22

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CC and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and

CC *Bordetella pertussis*, *malaria* plasmodia, *Leishmania*, *Trypanosoma* and *Schistosoma*. The immunomodulatory sequences are used to screen for human

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CC oligonucleotide; and determining the relative amount of Th1-biased

CC cytokines in the supernatant. Sequences AAV80104 to AAV80116 represent

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CC

SQ	Sequence 22 BP; 6 A; 4 C; 6 G; 6 T; 0 U; 0 Other;	Db	1 TGACTGTGAACTTGTAGATGA 22
Query Match	85.5%; Score 18.8; DB 2; Length 22;		
Best Local Similarity	90.9%; Pred. No. 24;		
Matches	20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;		
Qy	1 TGACTGTGAACTTGTAGATGA 22		
Db	1 TGACTGTGAACTTGTAGATGA 22		
RESULT 14			
AAV80104	Immunomodulatory oligo comprising an ISS sequence.		
ID	AAV80104 Standard; DNA; 22 BP.		
XX			
AAV80104;			
AC			
XX			
12-MAR-1999	(first entry)		
DT	12-MAR-1999 (first entry)		
DE	Oligo used in experiments for stimulation of cytokine production.		
XX			
KW	Immunomodulatory; immunostimulatory; octanucleotide; immune regulation;		
KW	ISS: cancer; allergy; asthma; hepatitis B infection; papillomavirus;		
KW	human immunodeficiency virus; influenza; herpes; M. tuberculosis; B. pertussis; malaria; plasmodia; Leishmania; Trypanosoma; Schistosoma.		
XX			
DE	DE		
XX			
KW	DE		
XX			
OS			
Synthetic.			
XX			
PN	WO9855495-A2.		
XX			
PD	10-DEC-1998.		
XX			
PP	05-JUN-1998; 98WO-US011578.		
XX			
PP	06-JUN-1997; 97US-0048793P.		
XX			
PR	(DYNNA-) DYNAVAX TECHNOLOGIES CORP.		
XX			
PA	Schwartz D, Roman M, Dina D;		
XX			
PA	PT		
XX			
PT	WPI; 1999-059898/05.		
XX			
DR	DR; 1999-059898/05.		
XX			
PT	Immunostimulatory oligonucleotides regulate the immune system - and		
PT	contain an immune-stimulating octanucleotide sequence; for treating		
PT	cancer, allergic and infectious diseases.		
XX			
PS	Example 1; Page 29; 63pp; English.		
XX			
PS	The invention relates to immunomodulatory oligonucleotides that comprise		
CC	at least 1 immunostimulatory octanucleotide sequence (ISS) where the ISS		
CC	sequences are selected from the group consisting of AGCTTCC, AGCTTCC,		
CC	GACGTTCC, and GACGTTCC. The immunomodulatory sequences are used to treat		
CC	patients needing immune regulation, such as those suffering from cancer,		
CC	an allergic disease and asthma. They are also used to prevent infectious		
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CC	and papillomavirus, Hemophilus influenza, Mycobacterium tuberculosis and		
CC	Bordetella pertussis, malarial plasmodia, Leishmania, Trypanosoma and		
CC	Schistosoma. The immunomodulatory sequences are used to screen for human		
CC	immunostimulatory activity by incubating macrophage cells and the		
CC	oligonucleotide; and determining the relative amount of Th1-biased		
CC	cytokines in the supernatant. Sequences AAV80104 to AAV80116 represent		
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CC	were found in experiments for the stimulation of cytokine production and		
CC	were found to lack immunostimulatory activity. The invention provides		
CC	specific claimed examples (AAV80096-103) of immunomodulatory sequences		
XX			
SQ	Sequence 22 BP; 7 A; 1 C; 8 G; 6 T; 0 U; 0 Other;		
Query Match	85.5%; Score 18.8; DB 2; Length 22;		
Best Local Similarity	90.9%; Pred. No. 24;		
Matches	20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;		
Qy	1 TGACTGTGAACTTGTAGATGA 22		
Db	1 TGACTGTGAACTTGTAGATGA 22		
Query Match	85.5%; Score 18.8; DB 2; Length 22;		
Best Local Similarity	90.9%; Pred. No. 24;		
Matches	20; Conservative 0; Mismatches 2; Indels 0; Gaps 0;		
Qy	1 TGACTGTGAACTTGTAGATGA 22		
Db	1 TGACTGTGAACTTGTAGATGA 22		

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